#### Commandant United States Coast Guard

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COMDTINST 6010.21A 1 NOV 1993

COMMANDANT INSTRUCTION 6010.21A

Subj: CLINICAL MONITORING AND EVALUATION (M&E)

Ref: (a) Medical Manual, COMDTINST M6000.1 (series)

- 1. <u>PURPOSE</u>. This instruction publishes additional Monitoring and Evaluation exercises for inclusion in Coast Guard health care facility Quality Assurance Programs.
- 2. ACTION. Area and district commanders; commanders, maintenance and logistics commands; commanding officers of Headquarters units; Commander, Coast Guard Activities Europe; and chiefs of offices and special staff divisions at Headquarters shall ensure compliance with the provisions of this notice.
- 3. <u>DIRECTIVES AFFECTED</u>. Commandant Instruction 6010.21, Clinical Monitoring and Evaluation (M&E), is canceled.
- 4. BACKGROUND. COMDTINST 6010.21, Clinical Monitoring and Evaluation (M&E), established the initial M&E schedule for Coast Guard clinics. This schedule was updated by COMDTNOTE 6010 of 8 JAN 93, CH-1 to COMDTINST 6010.21. The M&E schedule is further updated by this instruction. Section 13-H of reference (a) describes M&E schedule utilization.
- 5. <u>DISCUSSION</u>. Attached are two additional Medical M&E exercises (MED-3), two additional Dental M&E exercises (DENT-3), and two additional Drug Utilization Review M&E exercises (DUR-3).

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Clinics may use these exercises to supplement those previously provided. Administrative M&E exercises have been discontinued.

> /s/ ALAN M. STEINMAN Chief, Office of Health and Safety

- Encl: (1) FY 94 M&E Schedule and Clinical Aspects of Care for M&E
  - (2) M&E Exercises
  - (3) M&E Data Collection Log (4) M&E Flow Chart

### Non-standard Distribution:

- B:c MLCs (6 extra)
- C:a Cape Cod, Miami, Clearwater, Borinquen, Traverse City, Astoria only
- C:b North Bend, Port Angeles, Sitka only
- C:d Fort Macon, Miami Beach, Honolulu, Ketchikan only
- D:d Galveston and Humboldt Bay only

#### FY 94 MONITORING AND EVALUATION SCHEDULE

QUARTER	1st 	2nd 	3rd 	4th 
INITIAL	   MED-3 	DUR-3	   DENT-3	MED-4
*FOLLOW-UP	   DENT-2 	ADMIN-2	   MED-3 	DUR-3

### CLINICAL ASPECTS OF CARE FOR MONITORING AND EVALUATION

MED-1: Strep Throat

Urinary Tract Infection

MED-2: Gastroenteritis

Hypertension

MED-3: Nonspecific Vaginitis

Otitis Externa

DENT-1: Exodontia Informed Consent

Annual Dental Examinations

Dental Emergencies

DENT-2: Post-operative Infections

Restoration Replacements

DENT-3: Biopsies

Cast Restorations

DUR-1: NSAID Therapy

Antibiotic Therapy

DUR-2: Antihistamine Therapy

Antilipemic Therapy

DUR-3: Intranasal Steroid Therapy

Oral Contraceptive Therapy

### USING THE MONITORING AND EVALUATION SCHEDULE AND CLINICAL ASPECTS OF CARE LISTING

Each clinic shall initially monitor **one** clinic aspect of care each quarter. The schedule above determines the "menu" group for each quarter - the clinic selects an aspect of care from the listing for that group. For example, in the first quarter of FY94, M&E must be performed for an aspect of care on the medical (MED-3) menu (i.e., nonspecific vaginitis or otitis externa).

Completed M&E reports must be submitted to the Quality Assurance Focus Group (QAFG) prior to that last work day of each quarter. This means that data collection for each exercise should commence at the beginning of each quarter, in order to allow time for a representative data sample to be collected and evaluated prior to the end of the quarter. It is recommended that the QAFG assign responsibility for each exercise prior to the start of each quarter, so that data may be collected in a timely manner. Whenever possible, follow-up reports should be generated by the same person responsible for the initial M&E report.

#### \*FOLLOW-UP REPORTS

For studies that meet thresholds:

Each initial M&E Report must be followed 6 months later by follow-up reports.

For studies that do not meet thresholds:

A follow-up report is required 3 months after the initial report, and every 3 months after that, until thresholds are met.

Follow-up reports are recorded on the reverse side of the M&E Report form in sections 8, 9, and 10.

#### USING THE M&E DATA COLLECTION LOG (CG-5544)

Use this form, or a locally produced equivalent, while evaluating health records or other information sources for compliance with the indicator criteria. Record the indicator as being met, or not met, for each record reviewed. Indicate which indicator criteria, listed in Ssection 2 of the Monitoring and Evaluation Report, are not met by marking the appropriate column (e.g., (a), (b), etc.) on the log.

Retain completed logs, or equivalent, on file for three years for review by MLC QA site surveys.

1. Aspect of Care Diagnosis of acute minor illnesses: Nonspecific Vaginitis  2. Indicator All patients diagnosed with nonspecific vaginitis will have documentation in their health record of:  a. history of present illness, including sexual behavior; b. past medical history; c. medication history; d. temperature recorded; e. documented pelvic examination (positive or negative), including description of vaginal discharge; and f. microscopic examination of discharge to include: KOH prep (yeast), wet prep (clue cells) and cultures for chlamydia and GC.  (Five out of six criteria must be met.)  3. Threshold 90% of the records reviewed shall meet five or more criteria contained in the indicator.  4. Data Collection Methodology Use CLAMS and/or a review of patient records to retrospectively identify all patients with a diagnosis of "nonspecific vaginitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the MEE Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.  5. Evaluation Report Meet Date Evaluated:  7. Recommended Action  Signature Date  Oate	Facility	QA Coordinator				
a. history of present illness, including sexual behavior; b. past medical history; c. medication history; d. temperature recorded; e. documented pelvic examination (positive or negative), including description of vaginal discharge; and f. microscopic examination of discharge to include: KOH prep (yeast), wet prep (clue cells) and cultures for chlamydia and GC. (Five out of six criteria must be met.)  3. Threshold 90% of the records reviewed shall meet five or more criteria contained in the indicator.  4. Data Collection Methodology Use CLAMS and/or a review of patient records to retrospectively identify all patients with a diagnosis of "nonspecific vaginitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.  5. Evaluation Report  Meet Do Not Meet Indicator Criteria 6. Evaluator Name: Date Evaluated: 7. Recommended Action	1. Aspect of Care	Diagnosis of acute minor illnesses: Nonspecific Vaginitis				
b. past medical history; c. medication history; d. temperature recorded; e. documented pelvic examination (positive or negative), including description of vaginal discharge; and f. microscopic examination of discharge to include: KOH prep (yeast), wet prep (clue cells) and cultures for chlamydia and GC.  (Five out of six criteria must be met.)  3. Threshold 90% of the records reviewed shall meet five or more criteria contained in the indicator.  4. Data Collection Methodology Use CLAMS and/or a review of patient records to retrospectively identify all patients with a diagnosis of "nonspecific vaginitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.  5. Evaluation Report	2. Indicator All	patients diagnosed with nonspecific vaginitis will be documentation in their health record of:				
4. Data Collection Methodology  Use CLAMS and/or a review of patient records to retrospectively identify all patients with a diagnosis of "nonspecific vaginitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.  5. Evaluation Report  4 Meet  4 Do Not Meet Indicator Criteria  6. Evaluator  Name:  Date Evaluated:  7. Recommended Action	b. past medical hi c. medication hist d. temperature rec e. documented pelv description of f. microscopic exa wet prep (clue	<ul> <li>b. past medical history;</li> <li>c. medication history;</li> <li>d. temperature recorded;</li> <li>e. documented pelvic examination (positive or negative), including description of vaginal discharge; and</li> <li>f. microscopic examination of discharge to include: KOH prep (yeast), wet prep (clue cells) and cultures for chlamydia and GC.</li> </ul>				
records and/or a review of laboratory records to retrospectively identify all patients with a diagnosis of "nonspecific vaginitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.  5. Evaluation Report	3. Threshold 90% cri	of the records reviewed shall meet five or more teria contained in the indicator.				
6. Evaluator Name: Date Evaluated: 7. Recommended Action  Signature Date	records to retrospe "nonspecific vagini QAFG or its designe patients to determi have been met. Res each case, prior to not met, the QAFG s	records and/or a review of laboratory ctively identify all patients with a diagnosis of tis", up to a random sample size of 25 patients. The e will review the health record of all identified ne whether the criteria contained in the indicator ults may be logged on the M&E Data Collection Log for reporting results in section 5. If the threshold is hall review all cases which do not meet the criteria				
7. Recommended Action  Signature  Date	5. Evaluation Repor	t % Meet% Do Not Meet Indicator Criteria				
	6. Evaluator Name	Date Evaluated:				
	7. Recommended Acti	Signature Date				

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8. 3/6 month Follow-up Report  Continue M & E Recommended action:		<pre>% Meet the Evaluation % Do Not Meet Evaluat</pre>	Criteria ion Criteria M & E
		Signature	Date
9. 3/6 month Follow-up Report		* Meet the Evaluation * Do Not Meet Evaluation	Criteria ion Criteria
☐ Continue M & E Recommended action:		<pre>Discontinue</pre>	M & E
<del>,</del>	-	Signature	-/ <del></del>
10. 3/6 month Follow-up Report	J	% Meet the Evaluation % Do Not Meet Evaluat	Criteria
☐ Continue M & E			1 & E
Recommended action:			
		Signature	Date

Facility	QA Coordinator				
1. Aspect of Care	Diagnosis of Acute Minor Illnesses: Acute External Otitis (Otitis Externa)				
2. Indicator A	2. Indicator All patients diagnosed with Acute External Otitis will have documentation in their health record of:				
<ul><li>b. past medical</li><li>c. temperature r</li></ul>					
(2) descripti (3) descripti	of the tragus or auricle to elicit pain; on of the external ear canal; and on of any discharge found in the canal or external ear. or criteria must be met)				
3. Threshold 90	of the records reviewed shall meet at least three teria contained in the indicator.				
records to retrospectively identify all patients with a diagnosis of "acute external otitis", up to a random sample size of 25 patients. The QAFG or its designee will review the health record of all identified patients to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case, prior to reporting results in section 5. If the threshold is not met, the QAFG shall review all cases which do not meet the criteria prior to recommending improvement action in section 7.					
5. Evaluation Rep	rt   % Meet % Do Not Meet Indicator Criteria				
6. Evaluator Na	e: Date Evaluated:				
7. Recommended Ac	Signature Date				
	(continued on reverse )				

8. 3/6 month Follow-up Report  Continue M & E Recommended action:		* Meet the Evaluation * Do Not Meet Evaluat  Discontinue	ion Criteria
9. 3/6 month Follow-up Report		Signature  % Meet the Evaluation % Do Not Meet Evaluat:	Date Criteria ion Criteria M & E
Recommended action:			
10. 3/6 month Follow-up Report	\	Signature  * Meet the Evaluation	Date Criteria
☐ Continue M & E	<u> </u>	- % Do Not Meet Evaluat - Discontinue	de E
Recommended action:	_	Signature	/
		ordus cure	Date

Facility		QA Coordinator
1. Aspect of (	Care Bi	iopsies
2. Indicator	Biopsie	es are submitted properly:
and	_	erly prepared and shipped IAW USN SF-515 guidelines,
b. Adequate	tissue spe	ecimens are submitted.
3. Threshold sampling or in	forms I	than 10% of biopsy submissions shall have SF-515 returned with notes indicating inadequate tissue reparation and shipping.
4. Data Colle	ction Meth	hodology
a. Use CLAMS	or a log	to identify the number of biopsy procedures th, to a maximum of 10 cases.
b. Perform a the numbe	record re	eview of all biopsy cases (up to 10) to determine s returned with improper tissue samples or improper
preparati c. The QAFG	or its des	signee shall review the results to determine if the
threshold d. If the th and recom	reshold is	s exceeded, the QAFG shall review all biopsy cases on to the SDO.
5. Evaluation	Report	% Meet % Do Not Meet Indicator Criteria
6. Evaluator	Name:	Date Evaluated:
7. Recommende	d Action	
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		Signature / Date
		= (continued on reverse )

8. 3/6 month Follow-up Report  Continue M & E Recommended action:		% Meet the Evaluation % Do Not Meet Evaluation ① Discontinue	
	-	Signature	-/
9. 3/6 month Follow-up Report		<pre>% Meet the Evaluation % Do Not Meet Evaluat</pre>	Criteria ion Criteria
☐ Continue M & E Recommended action:		① Discontinue	
		Signature	-/ Date
10. 3/6 month Follow-up Report  Continue M & E  Recommended action:		* Meet the Evaluatio	tion Criteria M & E
		Signature	Date

Facility			QA Coordinator		
1. Aspect of (	Cast re	storations	3		
2. Indicator	Treatment is	planned p	properly;		
to prepara	to preparation of the teeth, and				
	<pre>(&lt; 6 months)</pre>	periapica r than 10%	tings shall have no record of recent all radiographs and periodontal of prepared teeth shall have no storation.		
4. Data Collec	tion Methodolo	<b>9</b> y			
month, to b. Perform a the number	a maximum of 1 record review	O cases. of all cas perative r	e number of castings placed in a sting cases (up to 10) to determine radiographs and probings or as.		
5. Evaluation	Report	ት Meet _	% Do Not Meet Indicator Criteria		
6. Evaluator	Name:		Date Evaluated:		
7. Recommended	Action				
	(co	ntinued or	Signature Date		

8. 3/6 month Follow-up Report  Continue M & E Recommended action:	Meet the Evaluation Criteria Do Not Meet Evaluation Criteria Discontinue M & E
	Signature / Date
9. 3/6 month Follow-up Report	* Meet the Evaluation Criteria * Do Not Meet Evaluation Criteria
☐ Continue M & E Recommended action:	Discontinue M & E
	Signature / Date
10. 3/6 month Follow-up Report  Continue M & E  Recommended action:	* Meet the Evaluation Criteria  Do Not Meet Evaluation Criteria Discontinue M & E
	Signature / Date

Facility		QA Coordinator	
1. Aspect of Ca	ste	g Utilization: Appropriate use of intranasal roids (Flunisolide, Beclomethasone, Dexamethasone, asonal and vasomotor rhinitis.	
2. Indicator	All pati	ents prescribed these medications will have:	
moderate to combined winder ventional to combination dosages and manufacture	severe so the a document of th	of seasonal or vasomotor rhinitis (characterized by symptoms lasting 4 weeks or longer per episode) mented history of unsuccessful treatment with concluding antihistamines, decongestants, or therapy, and many of administration within the product's proved guidelines, and that instruction in the proper use of these a provided.	
3. Threshold	95% of a criteria	all records reviewed will meet the indicator	
Use CLAMS or a review of prescription files to retrospectively identify all patients receiving prescriptions for intranasal steroids. The QAFG or its designee will review the health record to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case prior to reporting the results in section 5. If the threshold is not met, the QAFG or its designee shall review all cases which do not meet the criteria, evaluate, and recommend action in section 7.			
5. Evaluation 1	Report _	* Meet % Do Not Meet Indicator Criteria	
6. Evaluator	Name:	Date Evaluated:	
7. Recommended	Action	Signature Date	
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8. 3/6 month Follow-up Report		<pre>% Meet the Evaluation % Do Not Meet Evaluati</pre>	
Continue M & E Recommended action:			1 & E
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9. 3/6 month Follow-up Report		<pre>% Meet the Evaluation % Do Not Meet Evaluati</pre>	Criteria on Criteria
Continue M & E Recommended action:		☐ Discontinue	M & E
	,	Signature	-/ Date
10. 3/6 month Follow-up Report	V	% Meet the Evaluation	Criteria
☐ Continue M & E	J	No Not Meet Evaluat Discontinue	tion Criteria M & E
Recommended action:			
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		Signature	/
		princent	Date

Facility	QA Coordinator				
1. Aspect of Care	Drug Utilization: Appropriate use of oral contraceptive medications.				
2. Indicator All	2. Indicator All patients prescribed these medications will have:				
<ul> <li>a. documented history of physical exam within the past year including personal and family medical history, pelvic exam (including pap smear), breast exam, and vital signs, and</li> <li>b. documented evidence of instruction in the proper use of these medications including an explanation of side effects, missed doses, increased risk factors (smoking, etc.), and drug interactions (antibiotics, etc.).</li> </ul>					
••	of all records reviewed will meet the indicator teria.				
4. Data Collection Methodology  Use CLAMS or a review of prescription files to retrospectively identify all patients receiving prescriptions for oral contraceptives. the QAFG or its designee will review the health record to determine whether the criteria contained in the indicator have been met. Results may be logged on the M&E Data Collection Log for each case prior to reporting the results in section 5. If the threshold is not met, the QAFG or its designee shall review all cases which do not meet the criteria, evaluate, and recommend action in section 7.					
5. Evaluation Repor	t % Meet % Do Not Meet Indicator Criteria				
6. Evaluator Name	: Date Evaluated:				
7. Recommended Acti					
	Signature Date  — (continued on reverse )				

8. 3/6 month Follow-up Report  Continue M & E Recommended action:	% Meet the Evaluation % Do Not Meet Evaluati  Discontinue %	
	Signature	/ Date
9. 3/6 month Follow-up Report	% Meet the Evaluation % Do Not Meet Evaluati	
☐ Continue M & E Recommended action:	Discontinue 1	4 & E
	Signature	/
10. 3/6 month Follow-up Report  Continue M & E	* Meet the Evaluation * Do Not Meet Evaluat  Discontinue	
Recommended action:	<u> </u>	- <del>-</del> -
	Signature	/

# Enclosure (3) to COMDTINST 6010.21A M & E Data Collection Log

Facility	Aspect of Care
Data Collector	Date

	Date	Case Iduntification	Indicator Het Not Het	
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	Date	Case Identification	Indicator Met Not Met	
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### M & E FLOW CHART

